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## IN THE CLAIMS

Please amend the claims as follows. This listing of claims replaces all prior versions.

- 1-4. (Canceled).
- 5. (Previously presented) An isolated nucleic acid comprising a heterologous nucleotide sequence, a single retroviral long terminal repeat (LTR), a packaging signal, a rev responsive element, a polypurine tract, a eukaryotic promoter, a primer binding site, a bacterial origin of replication and a bacterial selection marker, and wherein the U3 region of the LTR comprises a loxP site.
- 6. (Previously presented) The nucleic acid of claim 5, further comprising a central polypurine tract.
- 7. (Previously presented) The nucleic acid of claim 5, further comprising a post-transcriptional regulatory element.
- 8. (Previously presented) A vector comprising the nucleic acid of claim 5.
- 9-11. (Canceled).
- 12. (Previously presented) The nucleic acid of claim 5, wherein the U3 region of the LTR comprises a restriction site.
- 13. (Previously presented) An isolated nucleic acid comprising a 5' retroviral LTR and a 3' retroviral LTR, a heterologous nucleotide sequence, a packaging signal, a rev responsive element, a polypurine tract, a enhanced promoter, a primer binding site, a bacterial origin of replication and a bacterial selection marker, wherein the bacterial origin of replication and bacterial selection marker are located between the two LTRs, and wherein the U3 region of the 3' LTR comprises a loxP site.

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- 14. (Previously presented) The nucleic acid of claim 13, further comprising a central polypurine tract.
- 15. (Previously presented) The nucleic acid of claim 13, further comprising a post-transcriptional regulatory element.

16-18. (Canceled).

- 19. (Previously presented) The nucleic acid of claim 13, wherein the U3 region of the LTR comprises a restriction site.
- 20. (Currently amended) A method of producing a single-LTR circular retroviral form plasmid, comprising:
- a. introducing a shuttle vector comprising the nucleic acid of claim 5 into a eukaryotic cell;
- b. extracting non-integrated DNA from the eukaryotic cell;
- c. transforming a bacterial cell with the DNA of step (b);
- d. selecting a bacterial cell showing expression of a selection marker; and isolating a single-LTR circular retroviral plasmid from the bacterial cell.
- 21. (Previously presented) A method of making a retroviral vector particle, comprising:
- a) introducing the vector of claim 8 into a retroviral packaging cell in medium, said packaging cell comprising nucleotide sequences encoding rev, gag/pol and env proteins but lacking packaging sequences; and
- b) collecting retroviral vector particles from the medium.
- 22. (Previously presented) A method of producing a retroviral expression vector, comprising cloning the nucleic acid of claim 5 into a non-retroviral expression vector.
- 23. (Previously presented) The retroviral expression vector produced by the method of claim 22.

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24-29. (Canceled).